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Listing of Claims

The following list of claims will replace all prior versions and listings of claims in the application.

1. (Currently Amended) A method of treating lymphoma in a subject in need thereof, comprising:

administering to a subject afflicted with lymphoma an antibody that binds to tenascin in a treatment effective amount, wherein said antibody is coupled to a radioisotope.

- 2. (Original) A method according to claim 1, wherein said antibody is a monoclonal antibody.
- 3. (Original) A method according to claim 1, wherein said subject is a human subject.
- 4. (Original) A method according to claim 1, wherein said antibody is selected from the group consisting of monoclonal antibody 81C6 and antibodies that bind to the epitope bound by monoclonal antibody 81C6.
- 5. (Original) A method according to claim 1, wherein said lymphoma is Hodgkin's lymphoma.
- 6. (Original) A method according to claim 1, wherein said lymphoma is Non-Hodgkin's lymphoma.

7. (Canceled)

8. (Currently Amended) A method according to claim 1 [[7]] wherein said radioisotope is selected from the group consisting of ¹³¹I, ⁹⁰Y, ²¹¹At, ²¹²Bi, ⁶⁷Cu, ¹⁸⁶Re, ¹⁸⁸Re, and ²¹²Pb.

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- 9. (Currently Amended) A method according to claim $\underline{1}$ [[7]], wherein said radioisotope is ^{131}I .
- 10. (Currently Amended) A method according to claim 1 [[7]], wherein said antibody coupled to a radioisotope is administered in an amount of from 5,000 rads to 100,000 rads 10 mCi to 100 mCi.
- 11. (Original) A method according to claim 1, wherein said administering step is a parenteral injection step.
- 12. (Original) A method of treating Non-Hodgkin's lymphoma in a human subject in need thereof, comprising:

parenterally administering to a human subject afflicted with lymphoma a monoclonal antibody that binds to tenascin in a treatment effective amount;

wherein said antibody is selected from the group consisting of monoclonal antibody 81C6 and antibodies that bind to the epitope bound by monoclonal antibody 81C6; and wherein said antibody is coupled to a radioisotope.

- 13. (Original) A method according to claim 12, wherein said Non-Hodgkin's Lymphoma is unresponsive to chemotherapy treatment selected from the group consisting of rituximab, cyclophosphamide, doxorubicin, vincristine and prednisone treatment.
- 14. (Original) A method according to claim 12, wherein said Non-Hodgkin's Lymphoma is unresponsive to rituximab treatment.
- 15. (Original) A method according to claim 12, wherein said radioisotope is selected from the group consisting of ¹³¹I, ⁹⁰Y, ²¹¹At, ²¹²Bi, ⁶⁷Cu, ¹⁸⁶Re, ¹⁸⁸Re, and ²¹²Pb.
 - 16. (Original) A method according to claim 12, wherein said radioisotope is ¹³¹I.

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17. (Original) A method according to claim 12, wherein said antibody coupled to a radioisotope is administered in an amount of from 5,000 rads to 100,000 rads 10 mCi to 100 mCi.

- 18. (Original) A method according to claim 12, wherein said Non-Hodgkin's lymphoma is a low grade lymphoma.
- 19. (Original) A method according to claim 12, wherein said Non-Hodgkin's lymphoma is an intermediate grade lymphoma.
- 20. (Original) A method according to claim 12, wherein said Non-Hodgkin's lymphoma is a high grade lymphoma.
- 21. (Original) A method according to claim 12, wherein said parenteral administration step is carried out by intravenous injection.
- 22. (New) A method of treating Non-Hodgkin's lymphoma in a human subject in need thereof, comprising:

parenterally administering to a human subject afflicted with Non-Hodgkin's lymphoma 81C6 monoclonal antibody coupled to ¹³¹I in an amount of from 10 mCi to 100 mCi, wherein the Non-Hodgkin's lymphoma is unresponsive to chemotherapy.